

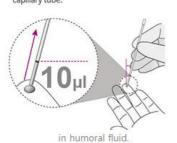
Instructions for use

TEST PROCEDURE - Be sure to test both STANDARD Q COVID-19 IqM and IqG simultaneously.

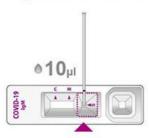
The test procedures for both COVID-19 IgM and IgG are the same.

Using Capillary whole blood

1 Collecting of Specimen Using a capillary tube, collect the 10µl of capillary whole blood to the black line of the capillary tube.



2 Adding of Specimen Add the collected capillary whole blood to the specimen well of the test device.



3 Dropping of buffer Add 3 drops (90µl) of buffer vertically into the buffer well of the test device.



4 Reading Time Read test result at 10-15 minutes.

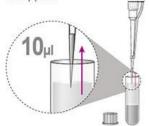




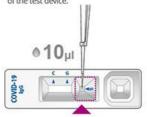
· Do not read test results after 15 minutes. It may give false results.

Using serum/plasma/venous whole blood

Collecting of Specimen Using a micropipette, collect the 10µl of serum, plasma or venous whole blood with micropipette.



2 Adding of Specimen Add the collected serum, plasma or venous whole blood to the specimen well of the test device.



3 Dropping of buffer

Add 3 drops (90µl) of buffer vertically into the buffer well of the test device.



4 Reading Time

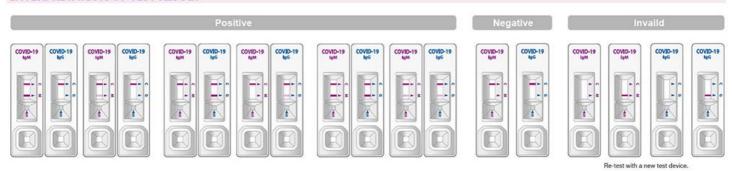
Read test result at 10-15 minutes.





· Do not read test results after 15 minutes. It may give false results.

INTERPRETATION OF TEST RESULT



- 1. A colored band will appear in the top section of the result window to show that the test is working properly. This band is control line (C).
- 2. A colored band will appear in the lower section of the result window. These bands are test line of IgM/IgG (M, G).
- 3. Even if the control line is faint, or the test line isn't uniform, the test should be considered to be performed properly and the test result should be interpreted as a positive result. * STANDARD Q COVID-19 IgM/IgG Duo Test may cross-react with antibody against SARS-CoV-1.
- * Results from antibody testing should not be used as the sole basis to diagnose or exclude SARS-CoV-2 infection or to inform infection status. * Positive results should be considered in conjunction with the clinical history, RT-PCR results and other data available.